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(54) Title: **DETERGENT OR PERSONAL CARE COMPOSITION WITH OIL CAPSULES**

(57) Abstract: Detergent or personal care compositions comprising capsules, which in turn comprise a hydrophobic continuous phase, the latter comprising an isotropic mixture of a thermoplastic-type co-polymer and hydrocarbon oil. Preferably, the capsules further comprise an oil-soluble benefit agent. Preferably, the compositions are liquid, especially aqueous-based compositions. In the preferred embodiment, the capsules are transparent or translucent.

DETERGENT OR PERSONAL CARE COMPOSITION  
WITH OIL CAPSULES

FIELD OF THE INVENTION

5 Detergent or personal care compositions containing capsules formed of a thermoplastic polymer and a hydrocarbon oil, and, preferably an oil-soluble benefit agent.

BACKGROUND OF THE INVENTION

10 In many articles of commerce, particularly consumer products, it is desirable to separate certain ingredients, yet have them disposed in a common container. Separation is particularly beneficial where one or more ingredients have negative interactions with each other. For example, in laundry  
15 detergents, enzymes are useful in removing stains but it is also best to separate them from other constituents, such as sources of alkalinity and surfactants, especially anionic surfactants like linear alkylbenzene sulfonates or alkyl sulfates. Bleaches, vitamins, perfumes, vegetable oils, plant  
20 extracts and ceramides are further examples of ingredients that sometimes need to be separated from the rest of the composition.

A known technique for separating ingredients in a common  
25 container includes encapsulation. Encapsulation technology is well known for different applications. Generally, encapsulation includes a medium that surrounds at least one component and thereby provides a barrier between the "encapsulated" component and other components. The barrier is typically temporary and  
30 is designed to break down and release the encapsulated material at a desired time, such as at a particular temperature, upon reaction or dissolution with chemicals, or due to mechanical stress. Methods of encapsulation include coacervation,

liposome formation, granulation, coating, emulsification, atomization and spray-cooling.

WO 92/20771 discloses particles having a substantially  
5 anhydrous core comprising a matrix polymer containing active ingredient, a layer of hydrophobic oil around the core and a shell of polymer around the oil layer. The process for making the particles includes the steps of dispersion in oil of particles of a matrix polymer containing active ingredient,  
10 dispersing this dispersion into the aqueous solution and causing a solid polymer shell to form around the droplets of the matrix particles.

Norbury et al. (U.S. Patent 4,976,961 and 5,013,473) disclose  
15 the addition of a polymer (which may be a polystyrene-polybutadiene-polystyrene block copolymer, e.g. Kraton® 1107) or a wax to an oil phase to increase the viscosity of the oil. The oil-polymer phase forms a core of the Norbury capsule; shell material is formed from another polymer.

20

Falholt et al. (U.S. Patent 4,906,396, UK 2 186 884, and EP 0 273 775) disclose a protective enzyme system wherein enzymes are dispersed in a hydrophobic material which is an organopolysiloxane oil or a high molecular weight hydrocarbon  
25 including a solid or waxy material. Especially preferred are said to be hydrophobic liquids which had been stabilized by suspending therein hydrophobic solid particles such as hydrophobic silica.

30 Mitchnick et al. (U.S. Patent 5,733,531) disclose a cosmetic sun-blocking composition containing particles which include a matrix and a UV-attenuating compound incorporated into the

matrix. A preferred encapsulating matrix comprises wax. Certain polymeric materials may also be employed.

Ratuiste et al. (U.S. Patent 5,589,370) discloses a process for  
5 encapsulation of sensitive materials, the capsule containing an oil dispersion holding an active and an outer polymer shell surrounding the oil dispersion. Tsaur et al. (U.S. Patent 5,441,660 and U.S. Patent 5,434,069) also disclose a capsule containing an oil dispersion containing an active and the  
10 polymer shell surrounding the dispersion. Another patent by Tsaur et al. (U.S. Patent 5,498,378) discloses wax capsules containing a structuring agent which may be a hydrophobic silica, a hydrocarbon material and organophilic clay. An example of a high molecular weight hydrocarbon is given as a  
15 hydrocarbon rubber or elastomers.

Morrison et al. (U.S. Patent 5,879,694) discloses transparent stiff gel candles comprising a hydrocarbon oil and a block co-polymer of a thermoplastic rubber (e.g., Kraton® series of  
20 polymers).

Despite numerous capsules in the prior art, a problem remains to produce a commercially attractive capsule which is stable—the encapsulated ingredient should not leach out of the capsule  
25 upon storage (especially important is the stability of capsules in liquid compositions)—but should release the protected ingredient with ease during normal use.

An additional challenge is that the capsules need to be  
30 manufactured with relative ease. For instance, in some prior art capsules the melt temperature of the encapsulating material may damage the encapsulated material during the encapsulation process. In the case of perfume encapsulation, for example,

many perfumes are essential oils which are volatile, and thus can particularly benefit from low temperature processing. Yet, typical encapsulating ingredients, e.g. wax, have a high melting temperature.

5

A further problem with capsules for consumer products is the manufacture of either transparent or colored capsules. The encapsulating material, e.g. wax, is typically opaque. In such capsules the opaque shell obscures the color; neither can the transparency be attained. Yet, it is frequently desirable to produce a transparent or colored capsule to increase the appeal of the consumer product.

#### SUMMARY OF THE INVENTION

15 The present invention includes detergent or personal care compositions comprising capsules, which in turn comprise a hydrophobic continuous phase, the latter comprising an isotropic mixture of a thermoplastic-type co-polymer and hydrocarbon oil. Preferably, the capsules further comprise an oil-soluble benefit agent. Preferably, the compositions are liquid, especially aqueous-based compositions. In the preferred embodiment, the capsules are transparent or translucent.

25 The following detailed description and the examples illustrate some of the effects of the inventive compositions. The invention and the claims, however, are not limited to the following description and examples.

#### 30 DETAILED DESCRIPTION OF THE INVENTION

Except in the operating and comparative examples, or where otherwise explicitly indicated, all numbers in this description indicating amounts of material or conditions of reaction,

physical properties of materials and/or use are to be understood as modified by the word "about." All amounts are by weight, unless otherwise specified.

5 For the avoidance of doubt the word "comprising" is intended to mean including but not necessarily "consisting of" or "composed of". In other words the listed steps or options need not be exhaustive.

10 The term "isotropic" as used herein means a mixture of a block co-polymer and a hydrocarbon oil which has minimum or no turbidity (i.e. light scattering) at 20°C.

The term "continuous" does not necessarily mean "isotropic".

The term "continuous" is used herein to denote the phase which  
15 is predominant in volume during emulsification or dispersion of discontinuous phase in the continuous phase.

The term "hydrocarbon oil" as used herein means a hydrocarbon oil having a maximum viscosity of about 10 kg/(m)(sec),

20 preferably no greater than about 5 kg/(m)(sec).

The term "wax" as used herein means a hydrophobic material which is a solid at 20°C. By "solid" is meant the ingredient is not mobile at 20°C.

25

The term "immiscible" as used herein means that discontinuous phase does not form a single phase with the continuous phase, i.e. the two phases form a partition between phases.

30 The term "transparent" as used herein includes both transparent and translucent and means that an ingredient, or a mixture, or a phase, or a capsule, or a composition, or a package according to the invention preferably has a transmittance of more than

25%, more preferably more than 30%, most preferably more than 40%, optimally more than 50% in the visible part of the spectrum (approximately 410-800 nm).

5 Alternatively, absorbency may be measured as less than 0.6 (approximately equivalent to 25% transmitting) or by having transmittance greater than 25% wherein % transmittance equals:  $1/10^{\text{absorbency}} \times 100\%$ . For purposes of the invention, as long as one wavelength in the visible light range has greater than 25%  
10 transmittance, it is considered to be transparent/translucent.

The capsules included in the inventive compositions comprise a hydrophobic continuous phase containing a benefit ingredient. The capsules may optionally include a further discontinuous  
15 phase.

#### CONTINUOUS PHASE

The continuous phase of the capsules included in the present invention comprises a hydrocarbon oil, a block co-polymer  
20 containing at least one rigid block and at least one flexible block, and, preferably, an oil-soluble benefit agent. The mixture of the hydrocarbon oil and the block co-polymer according to the present invention is isotropic at 20°C. It should be understood that since the co-polymer is not pourable  
25 at 20°C (indeed, it is solid), it may be difficult to combine the co-polymer with the oil at 20°C to ascertain whether the mixture is isotropic. According to the present invention, a mixture may be formed at any suitable temperature at which the liquefied co-polymer forms an isotropic liquid mixture with the  
30 oil. The copolymer/oil mixtures suitable for use in the present invention, however, remain isotropic after cooling to 20°C.

Preferably, especially in the case of colored capsules, the continuous phase is transparent and uncolored.

#### 5     Block Co-polymer

In one embodiment of the invention, the co-polymer employed in the continuous phase of the inventive capsules is selected from the group consisting of a triblock co-polymer, radial co-polymer, and multiblock co-polymer, the co-polymer comprising  
10 at least one triblock with a structure: rigid block---flexible block---rigid block. Preferably the rigid block is styrene-type polymer, and the flexible block is rubber-type polymer. By virtue of employing the rigid-flexible-rigid block co-polymer, the viscosity of the oil is increased, and the  
15 hardened continuous phase is formed, yet the resulting capsule is sufficiently soft and friable to release the discontinuous phase in normal use. The co-polymer blends uniformly with oil at a temperature which is much lower than the melting point of wax, thus allowing for encapsulation of temperature-sensitive  
20 ingredients, e.g. perfume, vitamins, vegetable oil, etc.

The preferred co-polymers are transparent and uncolored, in order to attain a transparent and uncolored continuous phase.

25 Examples of suitable co-polymers include but are not limited to those that are described in Morrison et al. (U.S. Patent 5,879,694) hereby incorporated by reference herein.

A triblock polymer, or a mixture of triblock polymer with  
30 diblock polymer, are preferred, since they may more easily form robust capsules, without the addition of a hydrophobic solid, yet resulting in capsules with a transparent shell.

Each of the triblock, radial block and/or multiblock copolymers in the invention contains at least two thermodynamically incompatible segments. By the expression thermodynamically incompatible with respect to the polymers, it is meant that the polymer contains at least two incompatible segments, for example at least one hard and one soft segment. In general, in a triblock polymer, the ratio of segments is one hard, one soft, one hard or an A-B-A copolymer. The multiblock and radial block copolymers can contain any combination of hard and soft segments, provided that there are both hard and soft characteristics. In the optional diblock copolymer, the blocks are sequential with respect to hard and soft segments.

Commercially available thermoplastic rubber type polymers which are especially useful in forming the compositions of the present invention are sold under the trademark Kraton® by Shell Chemical Company. The Kraton® rubber polymers are described as elastomers which have an unusual combination of high strength and low viscosity and a unique molecular structure of linear diblock, triblock and radial copolymers. Each molecule of the Kraton® rubber is said to consist of block segments of styrene monomer units and rubber monomer and/or comonomer units. Each block segment may consist of 100 or more monomer or comonomer units. The most common structure is the linear ABA block type; styrene-butadiene-styrene (SBS) and styrene-isoprene-styrene (SIS), which is the Kraton® D rubber series.

A second generation polymer of this general type is the Kraton® G series. This copolymer comprises a styrene-ethylene-butylene-styrene type (S-EB-S) structure. The Kraton® G series is preferred in the practice of the invention, as the copolymers of this series are hydrogenated and thus more thermally stable;

that is, decomposition is less likely to occur during blending of the G series polymers with the oil (the D series polymers having unsaturation within the rubber block). The Kraton® G rubbers are indicated as being compatible with paraffinic and naphthenic oils and the triblock copolymers are reported as taking up more than 20 times their weight in oil to make a product which can vary in consistency from a "Jello®" to a strong elastic rubbery material depending on the grade and concentration of the rubber.

10

The optionally blended diblock polymers include the AB type such as styrene-ethylenepropylene (S-EP) and styrene-ethylenebutylene (S-EB), styrene-butadiene (SB) and styrene-isoprene (SI).

15

The ABA structure of the Kraton® rubber molecule has polystyrene endblocks and elastomeric midblocks. This series of polymers is indicated as being a compounding ingredient or additive in adhesives, sealants and coatings, asphalt modification for roads and roofing, polymer modification, thermoset modification, and oil modification including use as viscosity index improvers, greases and gels. Certain grades of the Kraton® D series are also indicated as being useful as viscosity modifiers for formulating multi-grade motor oils.

25

International Patent Application No. W088/00603, published Jan. 28, 1988, by Francis et al. describes block copolymers which can be used as one or more components in the present invention.

These block copolymers are described as gels or gelloid liquid extended polymer compositions which can comprise an intimate mixture of a block copolymer containing relatively hard blocks and relatively elastomeric blocks. The additional polymer or copolymer material has at least partial compatibility with, and

30

a higher glass transition softening or melting temperature than, the hard blocks of the block copolymer. The copolymer additionally has at least 500 parts by weight of extender liquid per 100 parts of the block copolymer, the liquid being  
5 present to extend and soften the elastomeric blocks of the block copolymer. The extender liquid can be a hydrocarbon oil and/or a synthetic oil. The entire disclosure of this published application is incorporated herewith.

- 10 International Patent Application No. W088/00603 also refers to European Patent Application No. 224389 of Garmarra et al, published Jun. 3, 1987. This European patent application discloses styrene-diene block copolymer compositions and in particular discloses a mixture of triblock copolymers and a  
15 hydrocarbon oil, wherein the mixture of triblock copolymers comprises a triblock polymer having (a) styrene to ethylene-butylene ratio of 14 to 30 styrene blocks to 70 to 86 ethylene-butylene blocks, and (b) ethylene-butylene ratio of 31 to 35 styrene blocks to 65 to 69 ethylene-butylene blocks, and  
20 wherein the ratio of copolymer A to copolymer B is from about 15 to 85 to about 85 to 15. These compositions are said to be particularly useful as sealing materials. The disclosure of European Patent Application No. 224389 is also incorporated herein by reference.
- 25 U.S. Patent No. 5,221,534 discloses gels having a mineral oil and blends of di- and triblock copolymers. These gels are useful for health and beauty aid compositions. These health and beauty aid compositions contain at least two diblock or triblock copolymers and an effective amount of one or more  
30 cosmetic ingredient. Preferred compositions in U.S. Patent No. 5,221,534 contain both diblock and triblock copolymers, with the polymer blend being from about 5 to 95 wt % of diblock polymer to 95 to 5 wt % of triblock polymer. The entire

disclosure of U.S. Patent No. 5,221,534 is also incorporated herein by reference.

U.S. Patent No. 4,369,284 describes a transparent gel prepared from triblock copolymers and oils, including food and technical grade white petroleum mineral oils. The triblock copolymers therein give specific styrene end blocks to ethylene and butylene center blocks. The end block to ethylene and butylene center block ratio is given as being between 31:69 and 40:60. It is preferred under the present invention, however, that the end block to ethylene and butylene center block ratio be less than 31:69. The polymer content in the Examples of U.S. Patent No. 4,369,284 is from 5.9 to 25 percent. The disclosure of U.S. Patent No. 4,369,284 is incorporated herein in its entirety.

15

U.S. Patent No. 5,132,355 discloses polyethylene block copolymers of the A-B-A type, where in A is a "hard" block and B is a "soft" block. The gel made from this diblock copolymer is disclosed as being useful for complex molded candles when used with paraffin wax.

The preferred polymer is a triblock polymer of the Kraton® G type, in particular Kraton® G-1650. Kraton® G-1650 is an SEBS triblock copolymer which has a specific gravity of about 0.91, and is said to have a tensile strength of about 3.45 newton/m<sup>2</sup> as measured by ASTM method D-412-tensile jaw tester separation speed 25.4 cm/min. The styrene to rubber content of Kraton® G-1650 is said by the manufacturer to be about 29:71, and the Brookfield viscosity is about 8 kg/(m)(sec)(toluene solution, at 25°C, 25%w). The Shore A hardness is about 75.

For making the transparent capsules, preferably a mixture of Kraton® 1650 with Kraton® 1702 is employed, even though Kraton® 1650 is sufficient on its own. The mixture may be preferred in some cases, in order to increase the hardness of the capsules,  
5 while preserving transparency.

In a second embodiment of the invention, the diblock co-polymer may be employed (having rigid-flexible blocks), even in the absence of a triblock or radial co-polymer. Kraton® 1702 is a  
10 diblock co-polymer (styrene-ethylene/propylene). The properties of Kraton® 1702 make it more suitable for use as a viscosity modifier in making an emulsion. According to the second embodiment of the invention, when using Kraton® 1702, in the absence of other copolymers, a hydrophobic solid is added,  
15 in order to form capsules. Kraton® 1650, on the other hand, forms a gel, when mixed with oil. When using the mixture of two Kraton® polymers, the weight ratio of Kraton® 1650 to Kraton® 1702 is generally from 1:10 to 10:1, more preferably from 3:1 to 7:1, most preferably from 2:1 to 5:1, and optimally  
20 from 1:1 to 4:1.

The block co-polymer is employed in the inventive capsules generally in an amount of from 0.1% to 15%, more preferably from 0.5% to 10%, most preferably from 0.5% to 7%, and  
25 optimally from 1% to 4%, by weight of the continuous phase.

#### Hydrocarbon Oil

According to the present invention, any hydrocarbon oil may be employed, as long as: (1) it is an oil at room temperature  
30 (i.e., has a maximum viscosity of about about 10 kg/(m)(sec), preferably no greater than about 5 kg/(m)(sec)) and (2) it forms an isotropic mixture with the block co-polymer described above.

The isotropicity of the mixture may be determined by measuring light scattering (such as by turbidity measurement). If the turbidity is minimum or none at 20°C, the mixture is isotropic. Suitable apparatus for measuring turbidity is UV-visible

5 spectrophotometer (turbidity is measured in the visible light range), e.g. HP-8452: mixtures having a transmittance of at least 50%, preferably at least 70%, are suitable for use in the present invention.

10 Natural or synthetic hydrocarbon oil or mixtures thereof may be employed. Generally, the hydrocarbon oil may be a paraffinic oil, a naphthenic oil, vegetable oil, mineral oil or the like.

Examples include but are not limited to mineral oil, castor oil, vegetable oil, corn oil, sunflower oil, peanut oil, jojoba  
15 oil, 2-ethylhexyl oxystearate (and other alkyl oxystearates), acetylated lanolin alcohol, alkyl palmitates such as isopropyl palmitate, 2-ethylhexyl palmitate, glycerol triacetates, disopropyl adipate, dioctyl adipate (and other alkyl adipates), isopropyl myristate, C12 to C15 alcohol benzoates, and the  
20 like. As is evident from this list, some of these oils may also deliver a functional benefit, in addition to being a forming block of the capsules.

Most preferably, the oil is mineral oil, because it is both  
25 economic and most compatible with the block co-polymer.

Preferably, when Kraton® series block co-polymers are employed (i.e., styrene-elastomer block co-polymers), the oil is essentially free of silicone-containing oils, in order to  
30 obtain optimum isotropic mixtures. By "essentially free" is meant that in the Kraton®/oil continuous phase, the amount of silicone oil is preferably less than 2%, by weight of the

continuous phase, more preferably less than 1%, most preferably less than 0.5% and optimally is 0%.

The preferred oils are transparent and uncolored, in order to  
5 attain a transparent and uncolored continuous phase (although colored continuous phase is also included within the scope of the invention).

#### Optional Preferred Ingredients in the Continuous Phase

10 The continuous phase preferably includes an oil-soluble benefit agent, especially when the hydrocarbon oil that is chosen for forming the capsule, does not provide any functional benefit (as is, for instance the case with mineral oil).

15 This ingredient forms an isotropic mixture with co-polymer/hydrocarbon oil mixture when heated, but, on cooling, the mixture may or may not remain isotropic. Thus, the isotropicity of the copolymer/hydrocarbon oil (and thus the suitability of the chosen oil) should be tested in the absence  
20 of the oil-soluble benefit agent.

According to the present invention, by mixing the oil-soluble benefit agent with the block co-polymer/hydrocarbon oil mixture, its viscosity is increased. The increased viscosity  
25 is advantageous for several reasons: it entraps the benefit agent; it reduces the volatility of low boiling point benefit agents (e.g., perfumes); and it allows for increased deposition and/or increased substantivity of the benefit agent on the clothes or skin. Furthermore, with volatile or temperature-  
30 sensitive benefit agents (e.g., essential oils and perfumes), the capsule manufacture does not damage the benefit agent, due to the low melting point processing of block co-polymers

compared to the higher melting point of traditional encapsulating materials.

Suitable oil-soluble benefit agents include but are not limited to essential oils, perfumes, vitamins, vegetable oils, plant extracts, anti-wrinkle compounds, photoprotective agents, dye fixative agents, antioxidants, insecticides, soil repelling agents, soil release agents, anti-bacterial agents, cationic surfactants, lubricants, moisturizers.

The most preferred oil-soluble benefit agent is perfume, since perfumes are difficult to encapsulate at traditionally high encapsulating temperatures, and perfumes benefit from the increased viscosity, to optimize substantivity to the clothes and enhanced deposition.

15

Oil-soluble benefit agents are described in further detail below, under Detergent Compositions and Personal Care Compositions sections.

The continuous phase generally includes from 0% to 60%, more preferably from 10% to 50%, most preferably from 15% to 40%, and optimally from 20% to 35%, of the oil-soluble benefit agent in order to provide optimal benefits (% by weight of the total continuous phase).

25

A preferred ingredient for the continuous phase, in order to strengthen the capsules is a hydrophobic solid. This ingredient forms an isotropic mixture with co-polymer/oil mixture when heated, but, on cooling, the mixture will no longer be isotropic. Thus, the isotropicity of the copolymer/oil (and thus the suitability of the chosen oil) should be tested in the absence of the hydrophobic solid. Furthermore, if transparent capsules are desired, the addition

of a hydrophobic solid is not preferred. Examples of suitable hydrophobic solids include, but are not limited to wax, microcrystalline wax, fatty acid, hydrophobic silica, pigment (e.g., titanium dioxide), fatty alcohols, thermoplastic homo-  
5 polymers (preferably, polymers with melting point less than 95°C, to prevent boiling-out of the aqueous phase) such as polyethylene, polypropylene, and mixtures thereof.

Preferably, the hydrophobic solid is selected from paraffin  
10 wax, beeswax, micro-crystalline wax, polyethylene, polypropylene, most preferably paraffin wax or beeswax, due to their low price and easy processability.

The continuous phase generally includes from 0.1% to 60%, more  
15 preferably from 5% to 50%, most preferably from 10% to 40%, and optimally from 30% to 35% of the hydrophobic solid, in order to achieve the best balance between strength of the capsules and their friability in use (% by weight of the total continuous phase).

20

#### Optional Discontinuous Phase

The optional discontinuous phase of the capsules is itself and/or comprises a benefit agent and/or a colorant (which may be the same as or different from the benefit agent included in  
25 the continuous phase—for instance, the benefit agent in the discontinuous phase does not have to be oil-soluble). In some embodiments of the invention, the discontinuous phase is itself a benefit agent. In other embodiments, the discontinuous phase is itself a colorant (e.g. a solid pigment). Still in other  
30 embodiments the discontinuous phase serves as a vehicle for additional benefit agent and/or colorant. And still in other embodiments of the invention the discontinuous phase may itself be a benefit agent and/or colorant and also further include an

additional benefit agent and/or colorant. According to the present invention, the discontinuous phase is immiscible with the continuous phase. The discontinuous phase may be a solution (aqueous or oil), an oil, an emulsion, a dispersion or  
5 a solid. The preferred form of the discontinuous phase is an oil or a solution (oil or aqueous solution), due to the relative ease of incorporation of the oil or the solution into the continuous phase. The capsules may include more than one discontinuous phase.

10

An additional benefit agent may be present in an amount of from 0 to 100%, preferably 0.01 to 50%, more preferably 0.1 to 20%, by weight of the discontinuous phase.

15 Typical additional benefit agents include, but are not limited to a bleach, a bleach precursor, a surfactant, an enzyme, a whitening agent, a fabric softener, an anti-wrinkle compound, a dye fixative, dye transfer inhibitors, anti-redeposition polymers, soil release polymers, an anti-foam agent, a perfume,  
20 a silicone oil, a vegetable oil, a vitamin, a plant extract, a hydroxy acid, an anti-oxidant, an anti-bacterial agent, a moisturizer, and mixtures thereof.

The colorant may be a dye or a pigment. Dyes are preferable,  
25 since they are water-soluble and thus are more easily incorporated into capsules, compared to pigments which are typically not water-soluble. In one preferred embodiment of the invention, a water-soluble colorant is employed. Most preferably, a water-soluble dye is encapsulated, alone or in  
30 the mixture with a benefit agent, within a transparent, uncolored continuous phase.

If the additional benefit agent/colorant is oil-soluble, than an oil is chosen to carry the benefit agent/colorant in the discontinuous phase; if the benefit agent/colorant is water-soluble, than the discontinuous phase is an aqueous solution.  
5 Of course, as mentioned above, solids may be employed, without making a solution or a dispersion.

The discontinuous phase is present generally in an amount of from 0.01 to 45%, more preferably from 5 to 45%, most  
10 preferably from 10 to 40%, and optimally from 20 to 35% (% by volume of the capsule) in order to deliver sufficient benefit agent/colorant, provide adequate protection for the benefit agent/colorant and to maintain the ease of processing.

15 Most preferably, the capsule contains both the benefit agent and the colorant, within a transparent shell, to provide a visual signal to the consumer that a composition contains an additional beneficial ingredient:

20

#### Capsule Shape and Size

The preferred capsules are substantially spherical, particularly, when incorporated into transparent composition and/or transparent package, in order to provide a pleasing,  
25 commercially attractive appearance. Other shapes, however, such as star, disk, square and oval, are possible.

The size of the capsules is such as to render them suitable for incorporation into detergent or personal care compositions.  
30 Typical size range is from 300 $\mu$ m to 5,000 $\mu$ m, more preferably from 500 $\mu$ m to 3,000 $\mu$ m, most preferably from 800 $\mu$ m to 1,600 $\mu$ m, to provide visibility while ensuring uniform suspension.

## PROCESS OF MAKING THE CAPSULES

The inventive capsules may be prepared by any known encapsulation processes. Preferably, however, the capsules are  
5 prepared by the following process.

The preferred process comprises immersing droplets of an emulsion or a dispersion containing the continuous and discontinuous phases into an aqueous curing solution containing  
10 a high HLB surfactant and/or a super-wetting agent.

In the first step of the inventive process, an emulsion or dispersion is prepared by mixing the continuous and (optional) discontinuous phases, the latter being or containing the  
15 ingredient to be encapsulated, e.g. bleach solution or a vegetable oil. If no discontinuous phase is present, then solely the continuous phase ingredients are mixed. In the preferred embodiment, the co-polymer is melted, mixed with oil, then the oil-soluble benefit agent is mixed-in, then the  
20 optional discontinuous phase is added, with stirring (agitation), to ensure uniform mixing of the ingredients. The resulting solution/emulsion/dispersion is preferably kept at a temperature in the range from 40°C to 95°C. Most preferably, the use of direct heat is avoided. A most preferred  
25 temperature range is from 60°C to 75°C.

The resulting solution/emulsion/dispersion is directed, either as a stream, or dripping, into the curing solution containing a surfactant with a relatively high HLB value and/or a super-wetting agent, whereby the discrete capsules are formed.  
30 Optionally, pressure may be employed in ejecting the stream, in order to ensure that the stream penetrates the surface of the curing solution. The curing solution may also be chilled, stirred, and/or pressurized. The curing solution is prepared by

combining water and at least one surfactant with a high HLB value and/or a super-wetting agent.

The surfactants for the curing solution are selected from the group consisting of high HLB (7 to 25, preferably 10 to 20, most preferably 12 to 16) surfactants, preferably linear and branched nonionic such as Neodol® 25-12, 25-9 and Tergitol® 15-S-9. In the most preferred embodiment, the surfactant is Neodol® 25-12, which has a carbon chain length between 12 and 15, with 12 ethylene oxide groups per molecule.

The surface tension modifying agent or super-wetting agent is a highly efficient, low surface energy surfactant. Examples of super-wetting agents are as follows:

Super-wetting Agent	Supplier	Chemical Description
Zonyl®FSO	duPont	Fluoro chemical with ethylene glycol
Fluorad®	3MCompany	Fluorinated alkyl alkoxylate
DC Q2	Dow Corning	Polyoxy ethylene modified polydimethyl siloxane
Tergitol® 15-S	Union Carbide	Mixture of linear secondary alcohols reacted with ethyleneoxide
Surfynol® TG	Air Products & Chemicals	2,4,7,9-teramethyl-5-decyne-4,7-diol
Makon® OP-9	Stepan Chemical	Octyl phenol with 9 moles of ethylene oxide
Fluowet® OTN	Hoechst Celanese	Fluoroaliphatic oxyethylate
Forafac® 1157N	Atochem	Polyfluoroalkyl betaine
Silwet® L-77	OSI Specialty	Polyalkylene oxide modified heptamethyltrisiloxane

The most preferred super-wetting agent is Silwet® L-77 due to its ready availability and optimum performance.

The constituents in the curing solution are preferably present  
5 in the following ranges: water, 60% to 99%, most preferably 80% to 95%; surfactant and/or a super-wetting agent, 1% to 40%, most preferably 5% to 15% (all by weight of the curing solution).

10 In the preferred embodiment, the curing solution comprises both the high HLB surfactant and the super-wetting agent.

The super-wetting agent is preferably added as a pre-diluted solution by dripping along, as close to the capsule formation  
15 as possible, so that the super-wetting agent is on the surface of the curing solution.

The preferred curing solution contains a super-wetting agent generally in an amount of from 0.1 to 40%, more preferably from  
20 1 to 20%, most preferably from 2 to 10%, and optimally from 3 to 7% (% by weight of the curing solution).

The curing solution is preferably kept at a temperature in the range from 0°C to 50°C. The most preferred temperature range  
25 is from 10°C to 30°C.

In one preferred embodiment, the emulsion/dispersion of the continuous and discontinuous phases is caused to flow (preferably, under pressure) to form a stream which is directed  
30 into the curing solution. The stream breaks up into capsules within the curing solution. The stream can be defined by temperature, velocity, width and distance from the upper surface of the curing solution. The size of the orifice

through which the stream is directed and the pressure with which it is ejected will also affect the nature of the stream. In a preferred embodiment, the following operating parameters were found to produce capsules in the range of 200 $\mu$ m to 2500 $\mu$ m:  
5 emulsion temperature: 54-85°C; vessel pressure: 0-1.05 kg/cm<sup>2</sup>, most preferably 0.3-0.6 kg/cm<sup>2</sup>; nozzle distance from curing solution: 2.5-20cm, most preferably 17.5 cm; nozzle orifice diameter: 0.0125-0.25 cm; curing solution temperature: 0-50°C.

- 10 In an alternative preferred method for forming the capsules, the emulsion/dispersion is delivered to the curing solution by a plurality of nozzles: the emulsion is allowed to drip under the static head or the pressure. The dripping forms capsules upon contact with the curing solution. The size of the nozzle  
15 openings and the height of the liquid in vessel ("static head") containing the emulsion and the distance from the curing solution all play a part in the ultimate size of capsules.

In each embodiment, the curing solution is continually agitated  
20 during the emulsion addition, in order to distribute the formed capsules and keep the surface in motion.

In each of the above processes, the droplets/capsules advantageously have a density greater than that of the curing  
25 solution. As such, the formed capsules fall to the bottom of the receiving vessel and do not interfere with new droplets/capsules as they contact the surface of curing solution. Preferably, the density of the capsules is at least 1.0, most preferably from 1.01 to 2. Suitable methods for  
30 increasing the density include, but are not limited to the addition of solid inorganic material, (e.g. sugar or sorbitol), or any high density solute to the discontinuous phase of the emulsion.

Without being bound by theory, it is believed that the outer, hydrophobic surface of capsules attracts the hydrophobic portions of the surfactant molecules in the curing solution, thereby leaving the hydrophilic portions of the surfactant molecules extending from the outer surface of the capsules. Assuming this to be true, the capsules naturally repel each other due to hydrophilic molecule portions extending from the hydrophobic "shell". During processing this is advantageous because the capsules remain separate in solution.

In the final, optional, step of the process, in order to make the process continuous, the capsules may be collected out of the curing solution. If the capsules have density higher than that of the curing solution (i.e., higher than the density of water), then the vessel holding the curing solution may have one or more openings in the bottom of the vessel, for the capsules to drain out. The vessel volume may be continuously made up with a fresh curing solution.

20

If the capsule density is lower than that of water, then the capsules may be collected off the top of the solution by floating the top of the solution off into a collection vessel.

Again, the vessel volume may be continuously made up with a fresh curing solution. When this method of collection is employed, it is particularly important that the droplets penetrate the solution surface (e.g., the pressure is employed in forming droplets and/or a super-wetting agent is employed), so that the capsules are formed before the top layer of the curing solution is collected.

The capsules may be directly incorporated into a detergent or personal care composition, or the capsules may be stored in the form of a concentrated stock solution, which generally includes from 10 to 95% of water, more preferably from 20 to 80%, most preferably from 30 to 70%, and optimally from 35 to 60%, in order to attain the best balance between maximum concentration of the capsules and the stock's viscosity, the latter affecting its pumpability (% by weight of the total stock solution).

#### 10 DETERGENT COMPOSITIONS

Various detergent compositions include, but are not limited to laundry compositions, hard surface cleaners, dishwashing compositions. Detergent compositions may be in the form of powders, tablets, liquids, gels, water-soluble pouches and any combinations thereof.

Preferred detergent compositions are liquids or gels and, especially, aqueous-based compositions, since such compositions present a particular challenge, when benefit agents, e.g. oils or perfumes, need to be included. Thus, preferred liquid detergent compositions include from 10 to 80%, preferably from 30 to 70%, most preferably from 50 to 65% of water. Liquid detergent compositions generally have a pH from 6 to 13, preferably from 7 to 9.

25

The capsule concentration in the detergent composition is determined by the nature and/or amount of the benefit agent and/or colorant, and the size of the particles. Detergent compositions generally include from 0.01 to 20%, more preferably from 0.1 to 10%, most preferably from 0.2 to 5%, and optimally from 0.4 to 3%, of the inventive capsules (% by weight of the composition).

Preferred laundry compositions comprise, in addition to the capsules, a surfactant, in an amount from 1 to 70% more preferably from 10 to 50%, most preferably from 15 to 35%, and  
5 optimally from 15 to 30% (% by weight of the laundry composition). Suitable detergent and laundry surfactants are well known to one of ordinary skill in the art and may in general be chosen from anionic, nonionic, amphoteric, and cationic surfactants. Preferably, the surfactant in the  
10 laundry compositions is anionic and/or nonionic, especially linear alkylbenzene sulfonate, alkyl ether sulfate, alcohol ethoxylates and mixtures thereof.

In addition to the surfactant and the inventive capsules, the  
15 preferred laundry composition may include one or more well-known laundry ingredients, such as builders (from 0.1 to 40% for powders, from 0.1 to 20% for liquids), anti-redeposition agents, fluorescent dyes, perfumes, soil-release polymers, colorant, enzymes, etc.

20

Preferred detergent compositions according to the invention contain inventive capsules with encapsulated perfume in the continuous phase. Perfume concentration in such compositions is in general from 1% to 60%, more preferably from 5% to 50%, most  
25 preferably from 15% to 40% by weight of the capsules.

Most preferably, the capsules further contain bleach or bleach system, preferably in aqueous solution in the discontinuous phase. Any bleach suitable for detergent application may be  
30 included. Examples include, but are not limited to chlorine bleaches, peracids, bleach precursors, alone or with oxygen sources.

Bleach capsules for detergent compositions generally include from 0.5 to 15%, more preferably from 1% to 10%, most preferably from 3% to 8% and optimally from 4% to 7%, of a 5 bleach, in order to deliver an optimum benefit for minimum cost (% by weight of detergent composition).

Preferably, the detergent composition is a transparent composition containing transparent or colored capsules, and 10 packaged in the clear/transparent package.

#### PERSONAL CARE COMPOSITIONS

15 Personal care compositions according to the present invention may be solid or liquid, and include products which are rinsed off after application (e.g., shower gels, shampoos or soap bars) and product that are left on after application (e.g., cosmetic lotions, gels and creams). Various personal care 20 compositions include, but are not limited to soap bar compositions, facial or body cleansing compositions, shampoo compositions, conditioner compositions, and cosmetic compositions. Personal care compositions may be in the form of solution, lotion, cream, gel, solid and any combinations 25 thereof.

Preferred personal care compositions are liquids, gels and, especially, aqueous-based compositions, since such compositions present a particular challenge, when benefit agents, e.g. 30 vitamins or vegetable oils, need to be included. Thus, preferred liquid personal care compositions include from 0.1 to 99% preferably from 2 to 80%, most preferably from 10 to 60%,

of water. Liquid personal care compositions generally have a pH from 3 to 8, preferably from 5 to 7.

The capsule concentration in the personal care composition is determined by the nature and/or amount of the benefit agent and/or colorant. Personal care compositions generally include from 0.01 to 20%, more preferably from 0.1 to 10%, most preferably from 0.2 to 5% and optimally from 0.4 to 3%, of the inventive capsules (% by weight of the composition).

10

Preferred personal care compositions comprise, in addition to the inventive capsules, a cosmetically acceptable vehicle, in an amount from 0.1 to 70%, more preferably from 3 to 85%, most preferably from 5 to 95% and optimally from 10 to 99% (% by weight of the composition). Suitable vehicles are well known to one of ordinary skill in the art and may in general be chosen from isotropic liquid formulas or structured liquid formulas. Preferably, the vehicle in the personal care compositions is structured liquid formulas especially lamellar forming (structured) liquid formulas.

In addition to the vehicle and the capsules, the preferred personal care composition may include one or more well-known personal care ingredients, such as viscosity builders (from 0.1 to 30%), pH controllers (stabilizers) (from 0.005 to 20%).

Preferred personal care compositions are personal wash compositions, wherein the capsules contain a combination of a benefit agent and a colorant, with the oil-soluble benefit agent chosen from vitamins, antibacterial agents, vegetable oils, and mixtures thereof.

Vitamins include, but are not limited to vitamins A, E, C. Anti-bacterial agents include, but are not limited to Triclosan®. Vegetable oils include but are not limited to sunflower seed oil, safflower oil.

5

The personal wash compositions include, in addition to the capsules and the vehicle, a surfactant, especially Tegobetaine® (Cocamidopropyl Betaine). The surfactant is included generally in an amount from 0.1 to 10%, most preferably from 1 to 3%, and  
10 optimally from 2 to 6% (% by weight of the total composition).

Preferably, the personal care composition is a transparent composition containing transparent or colored capsules, and packaged in the clear/transparent package.

15

The following specific examples further illustrate the invention, but the invention is not limited thereto.

Suppliers and chemical description of the ingredients used in  
20 the examples are summarized in the following table:

Trade Name (if appropriate)	Chemical Name	Supplier
Kraton® 1650	Styrene-(ethylene-Butylene)-Styrene tri-block co-polymer	Shell
Kraton® 1702	Styrene-(Ethylene-Propylene) di-block co-polymer	Shell
Sorbitol	D-Glucitol	SPI Polyols
Neodol® 25-9	Nonionic surfactant: Alcohol Ethoxylate, C12-C15  9 Ethylene oxide groups HLB=13.1	Shell
Blue Dye	Sandolan Blue E-HRL powder 120	Sandoz Chemicals
Silwet® L-77	Polyalkylene oxide modified	OSI Specialties

	hertamethyltrisiloxane 84%, and Allyloxypolyethylene glycol methyl ether 16%	
	Mineral Oil	Fischer Scientific
1218® Wax	Petroleum, Hydrocarbon, paraffin Wax	IGI
Menthol Oil	Hexahydrothymol	Irvine Boody
	Vitamin E	BASF
Triclosan®	2,4,4 Trichloro- 2Hydroxydiphenyl Ether	Ciba
	Sodium Hypochlorite	Clorox
	Sunflower Seed Oil	Welch, Holmes and Clark Co., Inc.

## EXAMPLES 1-3

Several capsule formulations, within the scope of the invention, were prepared, using the apparatus and process as follows.

An encapsulation unit was built from stainless steel, jacketed, beaker. Nozzles were attached at the bottom of jacketed beaker, through which emulsion was sprayed to form encapsulates. At the top of the unit openings were made for pressurized air, a mixing shaft, and a pressure release valve.

Hoses were attached to upper and lower portions of beaker, through which water from a temperature controlled water bath flowed. This apparatus allowed for consistent indirect heat supply.

Kraton® 1650, which is a powdery substance, was converted to gel by mixing with mineral oil at 76.7°C. Wax was then added and a homogeneous isotropic liquid (continuous phase) was formed at 62.8°C. The emulsion was then prepared by emulsifying aqueous solution of sorbitol in the continuous

phase. The resulting emulsion was a very stable emulsion: it did not phase separate for 24 hours at 60°C. Capsules were produced using the re-circulating encapsulation unit described in the previous paragraph.

5

To simplify the preparation, a stock of 1.5% Kraton®1650/mineral oil mixture (named "Kraton® 1650 gel" in the Tables and Examples below) was prepared by mixing and heating the mixture of 1.5 parts of Kraton® 1650 and 98.5 parts of mineral oil at 76.7°C until it became an isotropic liquid.

Capsule formulations		
	Ingredient	Amount
Example 1		
Hydrophobic continuous phase	1.5% Kraton® 1650 gel	38.5%
	menthol oil	15.3%
	1218 wax	7.7%
Hydrophilic discontinuous phase	70 w/w% Sorbitol solution	38.4%
	Blue Dye	0.1%
Example 2		
Hydrophobic continuous phase	1.5% Kraton® 1650 gel	39%
	1218 wax	10%
	Vitamin E	15%
Hydrophilic discontinuous phase	70 w/w% Sorbitol solution	35.9%
	Blue Dye	0.1%
Example 3		
Hydrophobic continuous phase	1.5% Kraton® 1650 gel	39%
	1218 wax	10%
	Triclosan®	15%
Hydrophilic discontinuous phase	70 w/w % Sorbitol solution	35.9%

	Blue Dye	0.1%
--	----------	------

Examples 1-3 all resulted in discrete, stable, round-shaped capsules which did not stick together when they came in contact with one another. Capsules were stable (no leakage of 5 encapsulated ingredient) for at least 3 months at 20°C.

#### EXAMPLE 4

High content of sunflower seed oil in capsules is highly desired for personal care products. This example demonstrates 10 the preparation of transparent/translucent, high sunflower seed oil content capsules within the scope of the invention.

Capsule Formulation	
Kraton® 1650	5 g
Mineral oil	45 g
Sunflower Seed Oil	150 g
Curing solution	
Water	495.0 g
Neodol® 25-9	2.5 g
Silwet® L-77	2.5 g

15 Kraton® 1650 was mixed with mineral oil at about 82°C until the mix became transparent gel. Sunflower seed oil was then added and mixed-in until the mixture became isotropic, then the temperature was reduced to about 65°C. Capsules were prepared by dropping the emulsion by a pipette into curing solution, 20 collected by a net and stored in a jar with curing solution. These capsules were translucent, round, with the average size of 2500 µm, and did not agglomerate or stick together. The sunflower seed oil content in these capsules was 75%.

**EXAMPLE 5**

This example demonstrated the preparation of perfume capsules. Mineral oil, wax, Kraton® 1702 and oil-soluble perfume (Athena Extra, ex. Givaudan) were the main raw materials. 3 grams of Kraton® 1702 was added to 87 g of the mineral oil. The mixture was heated and stirred continuously until the Kraton® was completely dissolved in oil at 80°C. 10 g of wax was melted and added to this gel-like solution, to give the solution the extra strength. The mixture was stirred continuously to form an isotropic solution. The mixture was allowed to cool down to 41.6°C. After cooling, 10 g of perfume was added and stirred continuously to form a homogeneous solution. The mixture comprised 10% perfume. A pipette was used to dispense the resulting perfume mix into a curing solution bath. The curing solution consisted of 20% Neodol® 25-9 and 80% water. The capsules were about 2500 µm in size and the clarity was between transparent and translucent.

**EXAMPLES 6-7**

Two liquid detergent compositions, within the scope of the invention, containing inventive capsules are as follows:

Ingredients	Example 7 (g).	Example 8 (g).
Deionized water	301.68	301.68
50% NaOH solution	67.08	67.08
40% sodium xylene sulfonate	42.59	42.59
Linear Alkylbenzene Sulfonic acid	242.41	242.41
Neodol® 25-9	107.62	107.62
Coconut fatty acid	14.20	14.20
47.1% sodium silicate	81.63	81.63
40% Alcosperse® 725	8.52	8.52

Perfume Capsules from Example 5	5	10
Gum solution		
Gellan	0.45	0.31
Hydroxypropyl methylcellulose	0.00	0.45
Deionized water	90.05	89.74
Miscellaneous:	To 1000	To 1000
pH = 12.45		

## EXAMPLES 8-9

Two personal wash (liquid soap/shower gel) compositions, within the scope of the invention, containing inventive capsules are prepared as follows:

## EXAMPLE 8

Ingredients	% by wt.
Amphoteric (e.g., cocoamidopropyl betaine)	15-20%
Anionic surfactant (e.g., sodium laureth sulfate)	10-15%
<i>Sunflower Seed Oil Capsules of Example 14</i>	2
Humectants (e.g., glycerin)	1-3%
Cationic polymers (e.g., polyquaternium)	0.1-1.0%
Nonionic (e.g., PEG 80 Sorbitan monolaurate)	1-3%
Sodium Hydroxide	0.11
Opacifier (e.g., styrenet acrylate)	0.4
Preservative (e.g., DMDM hydantoin)	0.2
Fragrance	1.0
Water	to 100.0

## EXAMPLE 9

Ingredients	% by wt.
Sodium Lauroamphoacetate	7
Sodium Laureth Sulfate	14
Cetyl Acetate And Acetylated Lanolin Alcohol	0.5
Lauric Acid	2.5 - 3.0*
Sunflower Seed Oil	3
Vitamin E Capsules of Example 2	1
Cocamide Monoethanol amide	2
Glycerin	2
Guar Hydroxypropyl Trimonium Chloride	0.5
Citric Acid	1.2
Titanium Dioxide	0.2
DMDM Hydantoin/Iodopropynyl Butylcarbamate	0.2
EDTA	0.002
EHDP (Etidronic Acid)	0.02
Perfume	0.5
Water	to 100.0

**Claims**

1. A detergent composition comprising:
  - (A) from about 0.01% to about 20%, by weight of the composition of capsules comprising:
    - (a) a hydrophobic continuous phase comprising:
      - (a1) from about 0.1% to about 15%, by weight of the continuous phase, of a block co-polymer which is selected from the group consisting of a triblock co-polymer, radial co-polymer, and multiblock co-polymer, the co-polymer comprising at least one triblock with a structure: rigid block---flexible block---rigid block; and
      - (a2) a hydrocarbon oil which forms an isotropic mixture with the block co-polymer; and
    - (B) a surfactant.
2. The composition of claim 1 wherein the continuous phase further comprises from about 10% to about 60%, by weight of the continuous phase, of an oil-soluble benefit agent.
3. The detergent composition according to claim 1 wherein the continuous phase is transparent or translucent.
4. The detergent composition according to claim 1 wherein the composition is contained within the transparent package.
5. The detergent composition according to claim 1 wherein the composition is aqueous.
6. The detergent composition according to claim 1 wherein the capsule further comprises a colorant.

7. The composition according to claim 2 wherein the capsule further comprises from about 0.01% to about 45%, by volume of the capsule, of a discontinuous phase immiscible with the continuous phase.
8. The detergent composition according to claim 7 wherein the discontinuous phase is an aqueous solution comprising from about 1% to about 10%, by weight of the capsules, of a bleach.
9. A detergent composition comprising:
  - (A) from about 0.01% to about 20%, by weight of the composition of capsules comprising:
    - (a) a hydrophobic continuous phase comprising:
      - (a1) from about 0.1% to about 15%, by weight of the continuous phase, of a diblock co-polymer comprising at least one rigid block and at least one flexible block;
      - (a2) a hydrocarbon oil which forms an isotropic mixture with the block co-polymer;
      - (a3) from about 0.1% to about 60%, by weight of the continuous phase, of a hydrophobic solid; and
    - (B) a surfactant.
10. The detergent composition according to claim 9, wherein the continuous phase further comprises from about 10% to about 60%, by weight of the continuous phase of an oil-soluble benefit agent.
11. A personal care composition comprising:
  - (A) from about 0.01% to about 20%, by weight of the composition, of a capsule comprising:

- (a) a hydrophobic continuous phase comprising:
    - (a1) from about 0.1% to about 15%, by weight of the continuous phase, of a block co-polymer which is selected from the group consisting of a triblock co-polymer, radial co-polymer, and multiblock co-polymer, the co-polymer comprising at least one triblock with a structure: rigid block---flexible block---rigid block; and
    - (a2) a hydrocarbon oil which forms an isotropic mixture with the block co-polymer; and
  - (B) a cosmetically acceptable vehicle.
- 
- 12. The personal care composition according to claim 11, wherein the continuous phase further comprises from about 10% to about 60%, by weight of the continuous phase of an oil-soluble benefit agent.
  - 13. The personal care composition according to claim 11 wherein the continuous phase is transparent.
  - 14. The personal care composition of claim 11 wherein the composition is aqueous.
  - 15. The personal care composition according to claim 11 wherein the composition is contained within the transparent package.

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(57) Abstract: Detergent or personal care compositions comprising capsules, which in turn comprise a hydrophobic continuous phase, the latter comprising an isotropic mixture of a thermoplastic-type co-polymer and hydrocarbon oil. Preferably, the capsules further comprise an oil-soluble benefit agent. Preferably, the compositions are liquid, especially aqueous-based compositions. In the preferred embodiment, the capsules are transparent or translucent.

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Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	US 6 420 333 B1 (HSU ET AL) 16 July 2002 (2002-07-16) the whole document	1-3,5-7, 9-14
A	WO 92 20771 A (ALLIED COLLOIDS LTD) 26 November 1992 (1992-11-26) cited in the application claims; example 2	1,9
A	US 4 976 961 A (NORBURY ET AL) 11 December 1990 (1990-12-11) cited in the application claims; examples 1,4,5,10-13	11
A	US 5 013 473 A (NORBURY ET AL) 7 May 1991 (1991-05-07) cited in the application claims; examples	11
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<input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
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Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer  Serbetsoglou, A

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/EP 02/08641

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5 441 660 A (TSAUR ET AL) 15 August 1995 (1995-08-15) cited in the application column 3, line 18 -column 6, line 27; claims ---	1,8,9
A	US 5 434 069 A (TSAUR ET AL) 18 July 1995 (1995-07-18) cited in the application claims ---	1,8,9
A	US 5 498 378 A (TSAUR ET AL) 12 March 1996 (1996-03-12) cited in the application column 2, line 30 -column 3, line 50 column 5, line 41 -column 6, line 52 claims; examples ---	1,9
A	US 5 879 694 A (MORRISON ET AL) 9 March 1999 (1999-03-09) cited in the application column 3, line 1 -column 5, line 22; claims ---	1
A	US 5 221 534 A (DESLAURIERS ET AL) 22 June 1993 (1993-06-22) cited in the application claims; examples ---	1,11
A	WO 97 24179 A (ALLIED COLLOIDS LTD ;NOVO NORDISK A/S) 10 July 1997 (1997-07-10) claims; examples ---	1,9
A	US 5 972 508 A (BOECKH ET AL) 26 October 1999 (1999-10-26) claims 1-8,12 -----	1,8,9

## INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/EP 02/08641

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 6420333	B1	16-07-2002	NONE
WO 9220771	A	26-11-1992	US 5324445 A 28-06-1994 AT 143049 T 15-10-1996 AU 1789392 A 30-12-1992 CA 2102126 A1 15-11-1992 DE 69213934 D1 24-10-1996 DE 69213934 T2 30-01-1997 DK 585295 T3 17-03-1997 EP 0585295 A1 09-03-1994 ES 2093831 T3 01-01-1997 WO 9220771 A1 26-11-1992 GR 3021416 T3 31-01-1997 MX 9202246 A1 01-11-1992 US 5460817 A 24-10-1995 US 5492646 A 20-02-1996 ZA 9203510 A 17-05-1993
US 4976961	A	11-12-1990	AU 594789 B2 15-03-1990 AU 7464487 A 21-01-1988 BR 8703712 A 29-03-1988 CA 1295253 A1 04-02-1992 DE 3784803 D1 22-04-1993 DE 3784803 T2 12-08-1993 EP 0254447 A2 27-01-1988 KR 9609639 B1 23-07-1996
US 5013473	A	07-05-1991	NONE
US 5441660	A	15-08-1995	BR 9404433 A 20-06-1995 DE 69424506 D1 21-06-2000 DE 69424506 T2 14-09-2000 EP 0653485 A1 17-05-1995 ES 2147217 T3 01-09-2000
US 5434069	A	18-07-1995	BR 9404433 A 20-06-1995 DE 69424506 D1 21-06-2000 DE 69424506 T2 14-09-2000 EP 0653485 A1 17-05-1995 ES 2147217 T3 01-09-2000 ZA 9408975 A 13-05-1996
US 5498378	A	12-03-1996	NONE
US 5879694	A	09-03-1999	US 6066329 A 23-05-2000 AT 219134 T 15-06-2002 AU 708193 B2 29-07-1999 AU 7012296 A 19-03-1997 CA 2230312 A1 06-03-1997 DE 69621827 D1 18-07-2002 DE 871692 T1 31-08-2000 EP 0871692 A1 21-10-1998 ES 2179209 T3 16-01-2003 JP 11511498 T 05-10-1999 WO 9708282 A1 06-03-1997
US 5221534	A	22-06-1993	CA 2128423 A1 09-06-1994 WO 9412190 A1 09-06-1994

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 02/08641

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 5221534	A		DE 626855 T1	03-08-1995
			EP 0626855 A1	07-12-1994
			GR 95300037 T1	30-06-1995
			JP 2921986 B2	19-07-1999
			JP 7504441 T	18-05-1995
			AT 198703 T	15-02-2001
			DE 69231649 D1	22-02-2001
			DE 69231649 T2	03-05-2001
WO 9724179	A	10-07-1997	AT 216913 T	15-05-2002
			AU 716712 B2	02-03-2000
			AU 1203897 A	28-07-1997
			BR 9612314 A	13-07-1999
			CA 2239148 A1	10-07-1997
			CN 1208365 A	17-02-1999
			DE 69621045 D1	06-06-2002
			DE 69621045 T2	12-12-2002
			DK 876205 T3	12-08-2002
			EP 0876205 A1	11-11-1998
			ES 2174127 T3	01-11-2002
			WO 9724179 A1	10-07-1997
			JP 2000503052 T	14-03-2000
			NO 982999 A	26-08-1998
			PT 876205 T	30-09-2002
			US 6359031 B1	19-03-2002
			US 6225372 B1	01-05-2001
			ZA 9610879 A	23-12-1997
			AT 206073 T	15-10-2001
			AU 1203697 A	28-07-1997
			DE 69615582 D1	31-10-2001
			DE 69615582 T2	11-07-2002
			DK 873183 T3	21-01-2002
			EP 0873183 A1	28-10-1998
			ES 2164935 T3	01-03-2002
			WO 9724177 A1	10-07-1997
			JP 2000503051 T	14-03-2000
			PT 873183 T	28-03-2002
			US 6242405 B1	05-06-2001
US 5972508	A	26-10-1999	DE 19645024 A1	07-05-1998
			DE 59709000 D1	30-01-2003
			EP 0839902 A2	06-05-1998